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SIGNORINI, Raffaella [IT/IT]; Via Sanzio 3, I-35027 Noventa Padovana (IT).

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(74) Agent: APPOLONI, Romano; Notarbartolo & Gervasi S.p.A., Corso di Porta Vittoria, 9, I-20122 Milano (IT).

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(71) Applicant (for all designated States except US): UNIVERSITA' DEGLI STUDI DI MILANO - BICOCCA [IT/IT]; Piazza Ateneo Nuovo, 1, I-20126 Milano (IT).

(72) Inventors; and

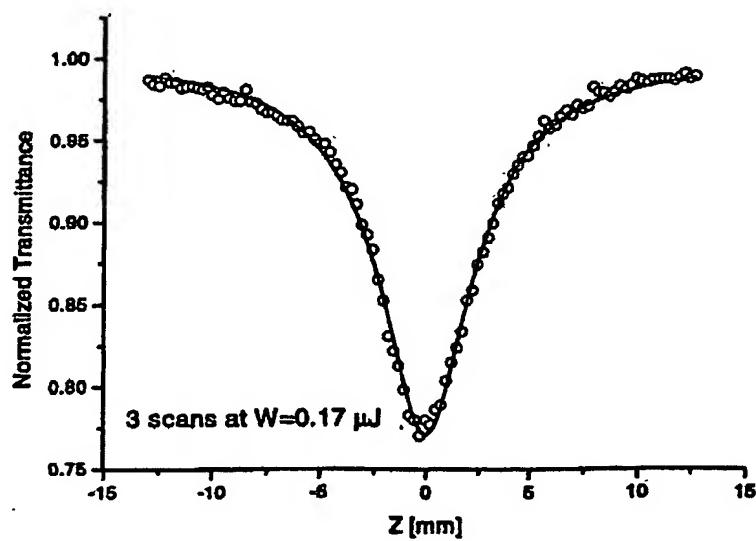
(75) Inventors/Applicants (for US only): ABBOTTO, Alessandro [IT/IT]; Via P. Caldirola, 6, I-20126 Milano (IT). BEVERINA, Luca [IT/IT]; Viale Corsica 72, I-20137 Milano (IT). BOZIO, Renato [IT/IT]; Via Piemonte, 11, I-35030 Selvazzano Destro (IT). PAGANI, Giorgio, A. [IT/IT]; Via Tarvisio, 19, I-20125 Milano (IT).

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(54) Title: TWO-PHOTON ABSORPTION HETEROAROMATIC CHROMOPHORES AND COMPOSITIONS THEREOF



WO 2004/018456 A1

(57) Abstract: The present invention relates to new heteroaromatic compounds with high two-photon absorption activity, useful in particular as optical power limiting agents via two-photon absorption or as imaging agents in confocal laser scanning fluorescence microscopy via two-photon absorption or excitation.



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Two-photon absorption heteroaromatic chromophores and compositions thereof

The present invention relates to new heteroaromatic chromophores with significant two-photon absorption activity.

5 It is known that molecules exhibit a nonlinear optical (NLO) behaviour by simultaneously absorbing two or more photons, either of the same or of different energy, to be promoted to one of their excited states when exposed to an intense laser pulse. In the case of two-photon absorption (TPA), the frequency of the nonlinear absorption is approximately half of that corresponding to the

10 conventional linear one-photon absorption. As a consequence, the TPA frequency typically falls in the visible red – near infrared (NIR) region of the electromagnetic radiation spectrum, where the material is transparent with respect to one-photon absorption. TPA is a 3rd order NLO process and is described by the imaginary part of the 3rd order nonlinear susceptibility.

15 Once the molecule has reached one of its excited states via TPA, it may show a fluorescence emission to return to its ground-state. In particular, the two-photon induced fluorescence emission occurs at a frequency very similar to the one-photon induced fluorescence emission. The direct consequence of this phenomenon is that the two-photon excited fluorescence frequency is usually

20 larger than the TPA frequency, as opposed to the case of linear absorption, where the emitted frequency is always smaller than that absorbed. Therefore, TPA dyes may absorb a red or NIR radiation (low frequency) and emit in the visible range.

Organic molecules able to show a significant TPA activity are very important for a variety of emerging applications including optical limiting (eye and sensor protection), three-dimensional optical memories, two-photon laser scanning fluorescence microscopy, up-converted lasing, non-destructive imaging of coated materials, and micro- and nanofabrication (MEMS, microelectromechanical systems) (Denk, W.; Strickler, J. H.; Webb, W. W. *Science* 1990, 248, 73; Ehrlich, J. E.; Wu, X. L.; Lee, L. Y. S.; Hu, Z. Y.; Rockel, H.; Marder, S. R.; Perry, J. W.

25 *Opt. Lett.* 1997, 22, 1843; Day, D.; Gu, M.; Smallridge, A. *Opt. Lett.* 1999, 24, 948; Cumpston, B. H.; Ananthavel, S. P.; Barlow, S.; Dyer, D. L.; Ehrlich, J. E.; Erskine, L. L.; Heikal, A. A.; Kuebler, S. M.; Lee, I. Y. S.; McCord-Maughon, D.; Qin, J. Q.;

Rockel, H.; Rumi, M.; Wu, X. L.; Marder, S. R.; Perry, J. W. *Nature* **1999**, *398*, 51; Belfield, K. D.; Ren, X. B.; Van Stryland, E. W.; Hagan, D. J.; Dubikovsky, V.; Miesak, E. J. *J. Am. Chem. Soc.* **2000**, *122*, 1217; Abbotto, A.; Beverina, L.; Bozio, R.; Bradamante, S.; Pagani, G. A.; Signorini, R. *Synth. Met.* **2001**, *121*, 1755; 5 Abbotto, A.; Beverina, L.; Bozio, R.; Bradamante, S.; Ferrante, C.; Pagani, G. A.; Signorini, R. *Adv. Mater.* **2000**, *12*, 1963).

The nonlinear absorption provides many advantages with respect to the conventional technologies based on linear absorption: a) two-photon excitation occurs in the red or NIR region; this region overlaps with the optical transparency 10 window of cells and living tissues; as a consequence, TPA provides much deeper light penetration depths as opposed to conventional techniques; b) the absorbed TPA intensity scales quadratically with the intensity I of the incident laser radiation, which in turn decreases approximately as the square of the distance from the 15 focus; the consequence is that molecules are excited via TPA only at the focus of the beam; two-photon induced phenomena occur only at the focus as well, with a fourth power increased spatial resolution; c) red and NIR light scattering is minimized with respect to higher frequency radiation.

The TPA phenomenon has been theoretically predicted by Göppert-Mayer in 1931. (Göppert-Mayer, M. *Ann. Phys.* **1931**, *9*, 273) and experimentally confirmed 30 years later (Kaiser, W. K.; Garrett, C. G. B. *Phys. Rev. Lett.* **1961**, *7*, 229). However, TPA has been studied in more detail only with the availability of proper laser sources. Moreover, all of TPA based applications remained unexplored for 20 decades due to the lack of efficient TPA absorbers. Only recently a number of dyes exhibiting significant TPA activity have been proposed. The vast majority of 25 these molecules are based on benzenoid derivatives substituted with conventional donor and acceptor groups such as NO_2 , CN , SO_nR , CO_2R , OR e NR_2 .

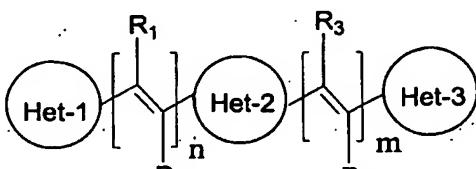
Few examples of TPA chromophores based on substituted heteroaromatic compounds are known. Concerning this aspect, WO 01/70735 owned by the same Applicant is mentioned.

30 In accordance with the present invention, new molecules are provided with high TPA activity, via excitation with lasers operating in the visible-red or NIR

wavelength region, that is a range where most organic molecular and polymeric materials and organic tissues are highly transparent.

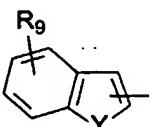
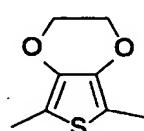
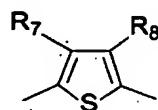
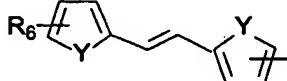
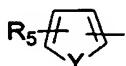
In accordance with the present invention, compounds are provided having the following general formulas (I) and (II)

5



(I)

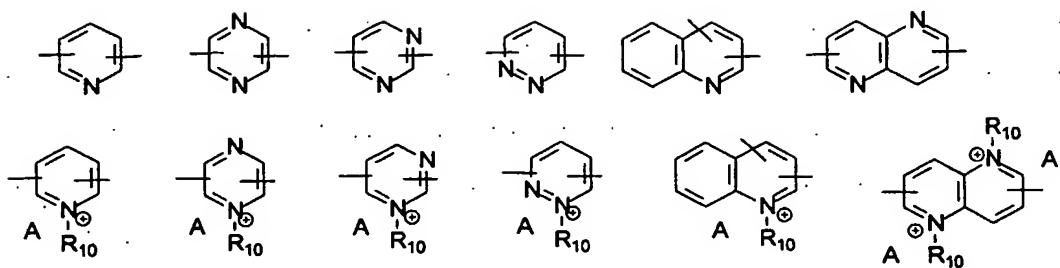
wherein Het-1 and Het-3 are identical or different, and are selected among the following heterocyclic groups:



10

wherein Y may be O, S, or NZ with Z = H, lower alkyl, and aryl; and wherein R₅, R₆, R₇, R₈, and R₉ are the same or different, and are selected from the group consisting of H, alkyl groups having from 1 to 18 carbon atoms, alkoxy, aminoalkyl, alkylhalide, hydroxyalkyl, alkyl groups containing hydroxy and amino functionalities, alkoxyalkyl, alkylsulfide, alkylthiol, alkylazide, alkylcarboxylic, alkylsulfonic, alkylisocyanate, alkylisothiocyanate, alkylalkene, alkylalkyne, aryl, formyl, and that can contain electronpoor ethenylic moieties such as maleimide, capable to react with nucleophilic groups such as -SH, and groups such as isothiocyanate capable to react with groups such as -NH₂;

15 and Het-2 is selected among the following heterocyclic groups:



wherein R_{10} is selected from the group consisting of H, alkyl groups having from 1 to 18 carbon atoms, alkoxy, aminoalkyl, alkylhalide, hydroxyalkyl, alkyl groups containing hydroxy and amino functionalities, alkoxyalkyl, alkylsulfide, alkylthiol,

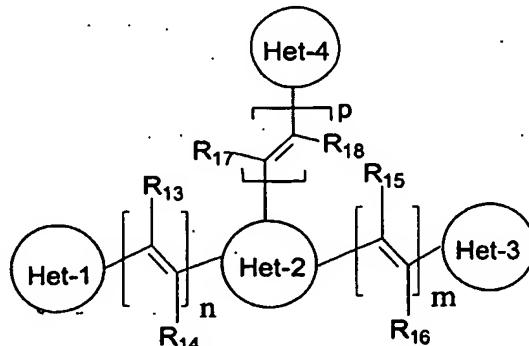
5 alkylazide, alkylcarboxylic, alkylsulfonic, alkylisocyanate, alkylisothiocyanate, alkylalkene, alkylalkyne, aryl, formyl, and that can contain electronpoor ethenylic moieties such as maleimide, capable to react with nucleophilic groups such as –SH, and groups such as isothiocyanate capable to react with groups such as –NH₂;

10 and A is selected among the anions alkylsulfonate, arylsulfonate, polyarenesulfonate, triflate, halide, sulfate, methosulfate, phosphate, polyphosphate;

and wherein n and m, the same or different may be 0,1,2;

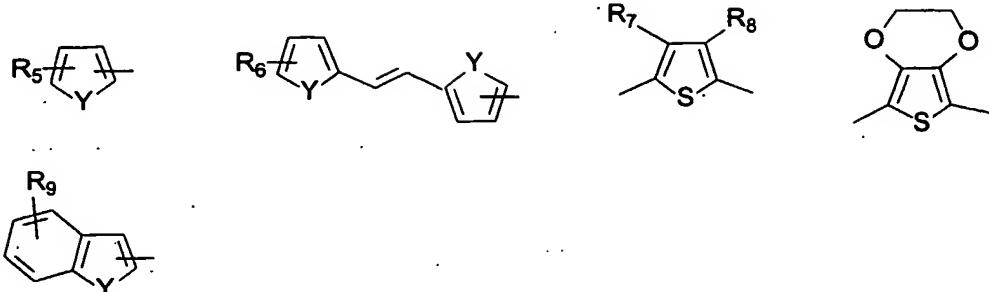
and R_1 , R_2 , R_3 , and R_4 , the same or different, may be H, lower alkyl, alkoxyalkyl,

15 aryl, cyano, alkoxy carbonyl, $-(CR_{11}R_{12})_p$ -Het, wherein $0 < p < 10$, R_{11} and R_{12} , the same or different, are selected from the group of H, lower alkyl, and Het may be Het-1 or Het-2 or Het-3.



(II)

20 wherein Het-1, Het-3, and Het-4 are the same or different and are selected among the following heterocyclic groups:

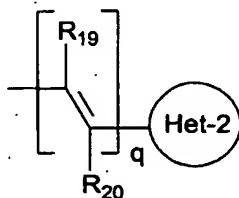


wherein Y may be O, S, and NZ with Z = H, lower alkyl, aryl;

and R₅, and R₆, are the same or different, and are selected from the group consisting of H, alkyl groups having from 1 to 18 carbon atoms, alkoxy, aminoalkyl,

5 alkylhalide, hydroxyalkyl, alkyl groups containing hydroxy and amino functionalities, alkoxyalkyl, alkylsulfide, alkylthiol, alkylazide, alkylcarboxylic, alkylsulfonic, alkylisocyanate, alkylisothiocyanate, alkylalkene, alkylalkyne, aryl, formyl, ketone, and that can contain electronpoor ethenylic moieties such as maleimide, capable to react with nucleophilic groups such as -SH, and groups

10 such as isothiocyanate capable to react with groups such as -NH₂; R₅, and R₆, the same or different, may further be the following heterocyclic group:



15 and R₇, R₈, and R₉ are defined as above;

and Het-2 is defined as above;

and wherein n, m, p, and q, the same or different, may be 0, 1, or 2;

and wherein R₁₃, R₁₄, R₁₅, R₁₆, R₁₇, R₁₈, R₁₉, and R₂₀ are the same or different and are selected from the group of H, lower alkyl, alkoxyalkyl, aryl, cyano,

20 alkoxycarbonyl, -(CR₂₁R₂₂)-Het, wherein 0<l<10, and R₂₁ e R₂₂, the same or different, are selected from the group of H, lower alkyl, and Het may be Het-1 or Het-2 or Het-3, or Het-4.

For the uses according to the present invention said compounds can show their two-photon absorption activity as such, or once prepared in solution, or in a solid state.

In a further aspect of the present invention said compounds can be processed into

5 compositions containing a polymer material such as poly(methacrylate), polyimide, polyamic acid, polystyrene, polycarbonate, and polyurethane or an organically-modified silica (SiO_2) network.

In particular, in said compositions the said compounds are either dispersed or covalently bonded to the polymer materials or to the silica network.

10 Features and advantages of the present invention will become readily apparent by reference to the following detailed description in conjunction with the accompanying drawings, in which:

Fig. 1 shows a typical two-photon absorption profile of compound (3) in DMSO (dimethylsulfoxide) obtained via the open-aperture Z-scan technique;

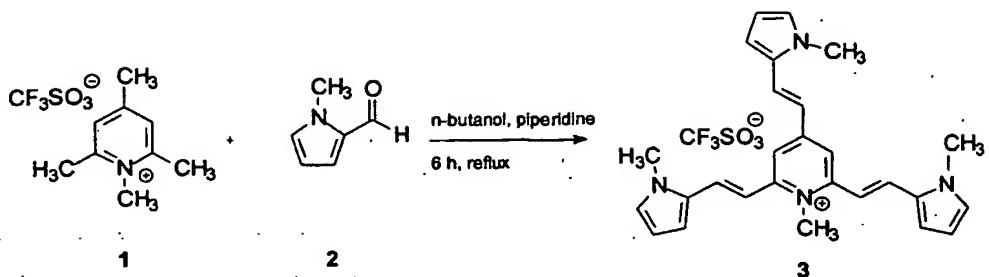
15 Fig. 2 shows a typical two-photon absorption profile of compound (6) in DMSO obtained with the same technique.

A detailed description of the invention is provided, with reference to certain compounds, which possess a structure corresponding to the formulas (3), (6), and (7), with examples which are not limiting the present invention.

20

EXAMPLE 1

Compound (3), endowed with TPA properties, has been prepared by a triple condensation reaction of compound (1) (Zhu, D.; Kochi, Jay K. *Organometallics* 1999, 18, 161) with an excess of *N*-methyl-2-pyrrolecarboxaldehyde in refluxing 25 *n*-butanol in the presence of a catalytic amount of piperidine as a base.

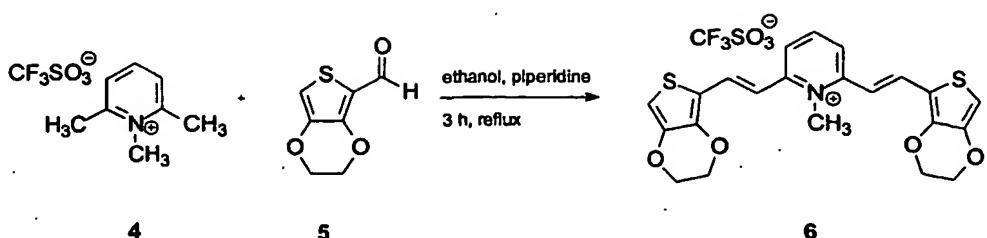


N-methyl-2,4,6-[1-(N-methylpyrrol-2-yl)ethen-2-yl]pyridinium triflate (3). A solution of *N*-methyl-2-pyrrolecarboxaldehyde (1.545 g, 14.16 mmol) in *n*-butanol (10 mL) was added dropwise to a solution of *N*-methyl-sym-collidinium triflate (1.332 g, 4.42 mmol) in the same solvent (40 mL). Ten drops of piperidine were added to the colorless solution and the mixture was stirred at reflux for 6 h. The resulting red-violet mixture was concentrated to ca. 15 mL and the red precipitate collected under reduced pressure. The solid was washed with toluene (10 mL) to give the product (1.783 g, 3.21 mmol, 68 %) mp 250 °C (dec) (*n*-BuOH); $^1\text{H-NMR}$ (CDCl_3) δ 7.78 (2 H, s), 7.63 (1 H, d, J = 16.0), 7.52 (2 H, d, J = 15.4), 6.90 (1 H, d, J = 16.0), 6.82 (1 H, d, J = 3.8), 6.78 (2 H, s), 6.75 (2 H, d, J = 15.4), 6.75 (2 H, d, J = 3.7), 6.72 (1 H, s), 6.21 (2 H, d, J = 3.2), 6.17 (1 H, d, J = 3.2), 3.95 (3 H, s), 3.83 (6 H, s), 3.82 (3 H, s); EA calcd for $\text{C}_{28}\text{H}_{29}\text{F}_3\text{N}_4\text{O}_3\text{S}$: C, 60.20 %; H, 5.23 %; N, 10.03 %. Found: C, 60.60 %; H, 5.29 %; N, 9.62 %.

We describe now a non limitative example related to a compound of general formula (I) and defined by the formula (6).

EXAMPLE 2

Compound (6) has been prepared by a condensation reaction of aldehyde (5) (Akoudad, S.; Frere, P.; Mercier, N.; Roncali, J. *J. Org. Chem.* 1999, 644267) and pyridinium salt (4) (Zhu, D.; Kochi, J. K. *Organometallics* 1999, 18, 161) in refluxing ethanol and in the presence of a catalytic amount of piperidine.

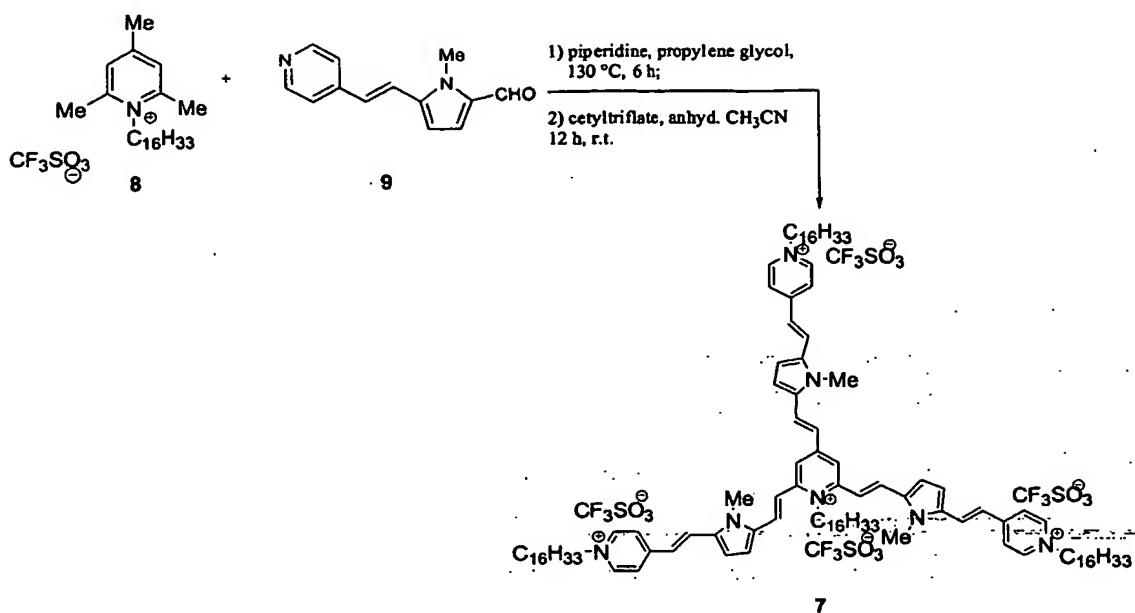


N-methyl-2,6-[1-(3,4-ethylenedioxythiophen-2-yl)ethen-2-yl]pyridinium triflate (6). A solution of 3,4-ethylenedioxythiophene-2-carboxaldehyde (0.456 g, 2.7 mmol) in ethanol (10 ml) was added dropwise to a solution of 1,2,6-trimethylpyridinium triflate (0.350 g, 1.3 mmol) and a few drops of piperidine in the same solvent (20 ml). Reaction mixture was refluxed for 3 hours and then cooled to 0 °C giving the formation of a brown-yellow precipitate that was filtered under reduced pressure and crystallized from ethanol (0.539 g, 0.94 mmol, 72%). mp 103-105 °C. $^1\text{H-NMR}$ (DMSO- d_6) δ 8.24 (1 H, t, J = 8.14), 8.16 (2 H, d, J = 8.09), 7.66 (2 H, d, J = 15.54), 7.11 (2 H, d, J = 15.63), 6.97 (2 H, s), 4.40 (4 H, m), 4.29 (4 H, m), 4.11 (3 H, s); $^{13}\text{C-NMR}$ (DMSO- d_6) 153.47 (2 C), 143.41 (2 C), 142.28 (1 C), 142.02 (2 C), 131.50 (2 C), 126.40 (2 C), 122.64 (2 C), 114.16 (2 C), 104.56 (2 C), 65.27 (2 C), 64.30 (2 C), 41.16 (2 C). Anal Calcd. for $\text{C}_{23}\text{H}_{20}\text{F}_3\text{NO}_7\text{S}_3$: C, 47.99 %; H, 3.50 %; N, 2.43 %. Found: C, 47.90 %; H, 3.11 %; N, 2.20 %.

We describe now another non limitative example related to a compound of general formula (II) and defined by the formula (7).

EXAMPLE 3

Compound (7) has been prepared following a two-step procedure. In the first step
 20 the *sim*-collidinium salt (8) has been condensed with an excess of aldehyde (9)
 (Abbotto, A.; Beverina, L.; Bozio, R.; Facchetti, A.; Ferrante, C.; Pagani, G. A.;
 Pedron, D.; Signorini, R. *Org. Lett.*, **2002**, *4*, 1495) in hot propylene glycol and in
 the presence of catalytic piperidine. Alkylation of the crude reaction product with
 an excess of cetyltriflate (Abbotto, A.; Bradamante, S.; Facchetti, A.; Pagani, G. A.
 25 *J. Org. Chem.* **1997**, *62*, 5755) in anhydrous acetonitrile gave the pure title
 compound.



N-cetyl-2,4,6-trimethylpyridinium triflate (8). A solution of cetyl triflate (1.873 g, 5 mmol) in dry toluene (5 ml) was added to a solution of *sym*-collidine (0.606 g, 5 mmol) under dry atmosphere. The white solution was heated at about 60 °C for 1 hour and then solvent was evaporated. The white residue was taken up with diethyl ether (10 ml) and filtered under reduced pressure, yielding the product as a white solid (1.982 g, 4 mmol, 80.0 %) mp 54-56 °C.

10 N-cetyl-2,4,6-[1-[N-methyl-5-(1-(pyrid-4-yl)-ethen-2-yl)pyrrol-2-yl]ethen-2-yl]pyridinium triflate (7). A solution of (9) (1.306 g, 6.1 mmol) in propylene glycol (15 mL) was added to a solution of *N*-cetyl-*sym*-collidinium triflate (0.460 mg, 0.93 mmol) and piperidine (5 drops) in the same solvent (10 mL). The resulting orange mixture was heated at 130 °C for 6 h yielding a dark violet solution which was cooled to room temperature and poured into Et₂O (100 mL). The obtained precipitate was collected by filtration under reduced pressure to give the monoquaternized precursor of (7) as a black solid, which was washed with water (20 mL) and EtOH (5 mL) (0.270 mg, 0.25 mmol, 27 %) mp >350 °C (dec); ¹H NMR (DMSO-*d*₆) δ 8.53 (6 H, d, J = 4.6), 8.23 (2 H, s), 7.99 (1 H, d, J = 15.7), 7.75 (2 H, d, J = 15.0), 7.62 (2 H, d, J = 16.2), 7.61 (1 H, d, J = 15.0), 7.60 (1 H, d, J = 14.8), 7.58 (6 H, d, J = 4.2), 7.26 (2 H, d, J = 15.2), 7.17 (2 H, d, J = 3.7), 7.10 (3 H, d, J = 15.9), 6.99 (1 H, d, J = 4.2), 6.91 (3 H, m), 4.60 (2 H, t, broad), 3.96 (3 H,

s), 3.95 (6 H, s), 1.78 (2 H, m, broad), 1.10-1.50 (26 H, m), 0.82 (3 H, t, J = 6.7); ^{13}C NMR (DMSO- d_6) δ 151.60 (2 C), 149.94 (6 C), 149.93 (1 C), 144.43 (3 C), 136.22 (1 C), 135.99 (2 C), 133.38 (1 C), 133.07 (2 C), 129.24 (2 C), 127.33 (1 C), 125.32 (1 C), 125.17 (2 C), 121.09 (3 C), 120.54 (6 C), 119.68 (1 C), 117.79 (2 C), 5 114.13 (2 C), 112.89 (2 C), 112.22 (1 C), 110.65 (1 C), 110.21 (2 C), 49.95 (1 C), 31.26 (1 C), 30.83 (1 C), 30.73 (2 C), 28.50-29.20 (12 C), 28.33 (1 C), 28.23 (1 C), 25.42 (1 C). A solution of cetyl triflate (2.050 g, 5.95 mmol) in anhyd. CH_3CN (40 mL) was added, under nitrogen atmosphere, to a solution of the product obtained as described in the previous step (1.195 g, 1.11 mmol) in the same solvent (80 mL). The reaction mixture was stirred overnight at room temperature and the solvent evaporated to leave a residue which was taken up with Et_2O (30 mL). The dark precipitate was collected by filtration under reduced pressure and washed several times with boiling hexane. The resulting blue solid was treated with boiling water to give the product (1.357 g, 0.62 mmol, 55.9 %). mp > 350 °C (EtOH); ^1H 10 NMR (DMSO- d_6) δ 8.80 (4 H, m), 8.63 (2 H, m), 8.30 (2 H, s), 8.26 (1 H, d, J = 14.1), 8.18-8.15 (4 H, m), 8.02-7.96 (3 H, m), 7.92-7.86 (2 H, m), 7.82-7.74 (3 H, m), 7.39 (1 H, d, J = 15.3), 7.31-7.24 (3 H, m), 7.23-7.14 (3 H, m), 7.12-7.09 (2 H, m), 7.05 (1 H, m), 7.01 (1 H, m), 4.68 (2 H, t broad), 4.45 (6 H, t broad), 3.98 (6 H, s), 3.97 (3 H, s), 1.90 (6 H, m broad), 1.77 (2 H, s broad), 1.45-1.10 (104 H, m), 15 20 0.90-0.80 (12 H, m).

According to the present invention said compounds show large two-photon absorption cross-sections both in solution and in the solid state. We now describe, with examples which are not limiting the present invention, experimental data of the two-photon absorption activity of said compounds (3), (6), and (7).

25 We define the following parameters: β (two-photon absorption coefficient, concentration dependent), σ_2 and σ_2' (cross-sections). It is possible to obtain the absorption coefficient β by interpolation of the relationship between the transmittance T versus the initial laser beam intensity I_0 , in accordance with the following relationships:

30

$$T = \frac{\ln(1 + I_0 L \beta)}{I_0 L \beta}$$

where $T = \frac{I_t}{I_0}$ and $L=1$ cm and I_t is the intensity of the transmitted beam.

The I_0 and I_t dimensions are $I_t, I_0 = [GW/cm^2]$ whereas the β dimensions are $\beta = [cm/GW]$

Since $\sigma_2 = \frac{\beta}{N_a}$ it follows that $\sigma_2 = \frac{\beta}{N_a c} 10^3$ where N_a is the Avogadro's number

5 and σ_2 has the dimensions of $[cm^4/GW]$.

Finally: $\sigma'_2 = h\nu \sigma_2$. σ'_2 has the dimensions of $\left[\frac{cm^4 \cdot s}{photon \cdot molecule} \right]$.

The following-table summarizes the nonlinear optical characterization data for said compounds taken as examples.

Compound	λ (nm)	Pulse duration (fs)	Power (μ J)	Intensity (GW/cm^2)	Concentration (mmol/l)	β (cm/GW)	σ'_2 $\left[\frac{10^{-50} cm^4 \cdot s}{photon \cdot molecule} \right]$
3	785	130-150	0.14	100	29.0	0.078	113
6	785	130-150	0.22	228	30.4	0.027	37
7	800	150			2.1		1600

10

Figures 1 and 2 show, as an example, the two-photon absorption activity of compounds (3) and (6), respectively. The TPA activity has been characterized by means of "open-aperture" Z-scan measurements of DMSO solutions of the described compounds and with a laser source operating at 780-790 nm with a pulse duration of 130-150 fs.

15

The Z-scan technique is one of the two most important experimental procedures to measure nonlinear absorption phenomena (two-photon absorption). The open-aperture Z-scan enables the measurement of the nonlinear absorption of the sample by recording the transmittance T (the ratio between transmitted and incident intensity) as a function of the incident intensity. To do this, the sample is moved along the propagation direction (the Z axis) of a focused laser beam. The energy of the laser beam is kept constant, while the intensity grows up as the sample moves towards the focal plane ($Z = 0$). Only the linear transmittance

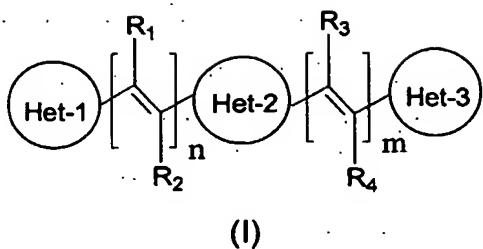
contributes to the signal far from the focal plane. In the proximity of the focus the intensity grows up very quickly and the nonlinear absorption process generates a dip in the transmittance ($T < 1$). The dip is symmetrical with respect to the position of the focal plane. When a fs source is employed, the Z-scan allows for the 5 discrimination between simultaneous TPA and sequential multiphoton absorption processes, involving intermediate excited states populated by nonradiative phenomena. In fact, in the case of fast (100 - 200 fs) pulses, the latter process does not contribute to the signal, being the nonradiative processes active in the picosecond (ps) or nanosecond (ns) regime. Since the two-photon absorption 10 scales quadratically, and not linearly, with the intensity of the incident laser radiation, the measured absorption as a function of the incident intensity provides unequivocal evidence that the sample is a non-linear (two-photon) absorber. In this way the two-photon absorption parameters β and σ_2' are experimentally obtained.

15 Figures 1 and 2 show the Z-scan profiles for DMSO solutions of molecules (3) and (6), respectively, measured in a 1-mm cell and with pulse energy of 0.17 and 0.16 mJ. The normalized transmittance ($I(z)/I(\infty)$, where $I(\infty)$ is the transmitted intensity far from the focal plane) is plotted as a function of the sample position (Z). The deep dip shown in both graphs is a clear evidence that a strong two-photon 20 absorption is occurring in solution. In addition, the Figures ensure that the two compounds show no significant linear absorption at 785 nm and, therefore, are completely transparent at low intensities of the incident radiation (Z far from the focal plane). Figure 1 proves that compound (3) shows a transmittance $T = 0.77$ at the focal point with a laser pulse energy of 0.17 mJ. This value is remarkably 25 lower, that is the two-photon absorption is larger, than that obtained for other known molecules of comparable molecular weight using the same laser power and pulse width conditions.

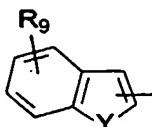
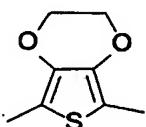
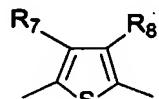
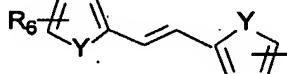
In a further aspect of the present invention, in addition to optical limiting activity, said compounds are useful for other applications based on their two-photon 30 absorption activity, such as use as imaging agents in confocal laser scanning fluorescence microscopy via two-photon absorption or excitation.

CLAIMS

1. A compound of formula (I).



wherein Het-1 and Het-3 are identical or different, and are selected among the following heterocyclic groups:



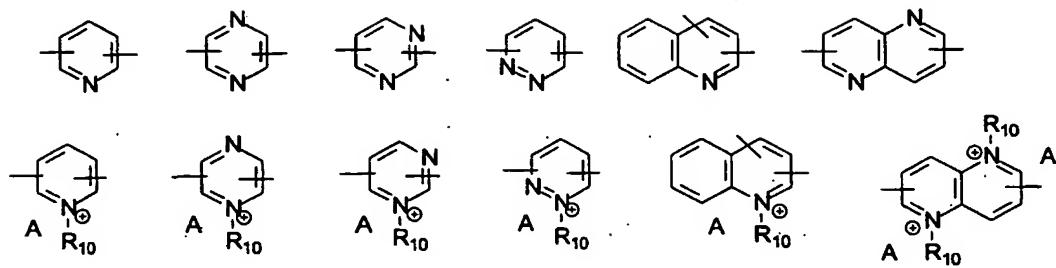
10

wherein Y may be O, S, or NZ with Z = H, lower alkyl, and aryl; and wherein R₅, R₆, R₇, R₈, and R₉ are the same or different, and are selected from the group consisting of H, alkyl groups having from 1 to 18 carbon atoms, alkoxy, aminoalkyl, alkylhalide, hydroxyalkyl, alkyl groups containing hydroxy and amino functionalities, alkoxyalkyl, alkylsulfide, alkylthiol, alkylazide, alkylcarboxylic, alkylsulfonic, alkylisocyanate, alkylisothiocyanate, alkylalkene, alkylalkyne, aryl, formyl, and that can contain electronpoor ethenylic moieties such as maleimide, capable to react with nucleophilic groups such as -SH, and groups such as isothiocyanate capable to react with groups such as -NH₂;

15

20

and Het-2 is selected among the following heterocyclic groups:



wherein R₁₀ is selected from the group consisting of H, alkyl groups having from 1 to 18 carbon atoms, alkoxy, aminoalkyl, alkylhalide, hydroxyalkyl, alkyl groups containing hydroxy and amino functionalities, alkoxyalkyl, alkylsulfide, alkylthiol,

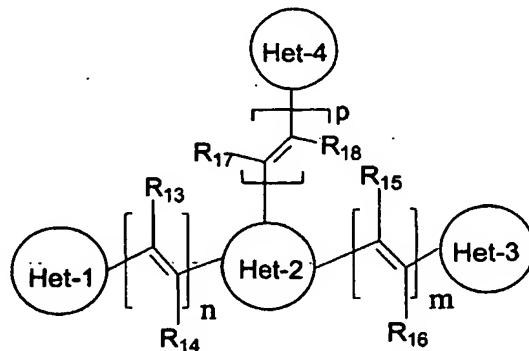
5 alkylazide, alkylcarboxylic, alkylsulfonic, alkylisocyanate, alkylisothiocyanate, alkylalkene, alkylalkyne, aryl, formyl, and that can contain electronpoor ethenylc moieties such as maleimide, capable to react with nucleophilic groups such as – SH, and groups such as isothiocyanate capable to react with groups such as – NH₂;

10 and A is selected among the anions alkylsulfonate, arylsulfonate, polyarenesulfonate, triflate, halide, sulfate, methosulfate, phosphate, polyphosphate;

and wherein n and m, the same or different may be 0,1,2;

and R₁, R₂, R₃, and R₄, the same or different, may be H, lower alkyl, alkoxyalkyl, 15 aryl, cyano, alkoxy carbonyl, -(CR₁₁R₁₂)_p-Het, wherein 0<p<10, R₁₁ and R₁₂, the same or different, are selected from the group of H, lower alkyl, and Het may be Het-1 or Het-2 or Het-3.

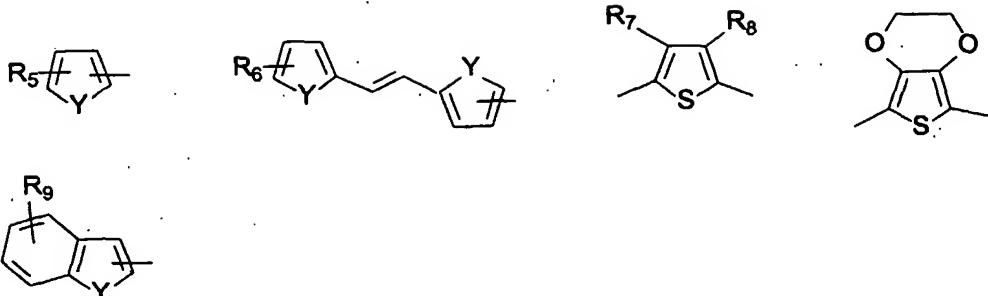
2. A compound of formula (II)



(II)

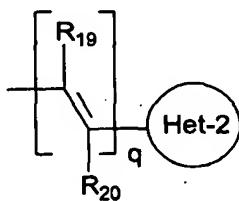
20

wherein Het-1, Het-3, and Het-4 are the same or different and are selected among the following heterocyclic groups:



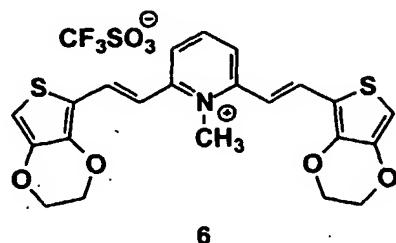
wherein Y may be O, S, and NZ with Z = H, lower alkyl, aryl;

5 and R₅, and R₆, are the same or different, and are selected from the group consisting of H, alkyl groups having from 1 to 18 carbon atoms, alkoxy, aminoalkyl, alkylhalide, hydroxyalkyl, alkyl groups containing hydroxy and amino functionalities, alkoxyalkyl, alkylsulfide, alkylthiol, alkylazide, alkylcarboxylic, alkylsulfonic, alkylisocyanate, alkylisothiocyanate, alkylalkene, alkylalkyne, aryl, formyl, ketone, and that can contain electronpoor ethenylic moieties such as maleimide, capable to react with nucleophilic groups such as -SH, and groups such as isothiocyanate capable to react with groups such as -NH₂; R₅, and R₆, the same or different, may further be the following heterocyclic group:

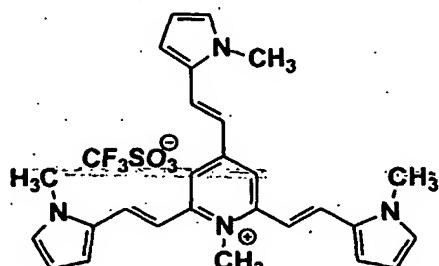


15 and R₇, R₈, and R₉ are defined as in claim 1;
 and Het-2 is defined as in claim 1;
 and wherein n, m, p, and q, the same or different, may be 0, 1, or 2;
 and wherein R₁₃, R₁₄, R₁₅, R₁₆, R₁₇, R₁₈, R₁₉, and R₂₀ are the same or different and are selected from the group of H, lower alkyl, alkoxyalkyl, aryl, cyano, alkoxycarbonyl, -(CR₂₁R₂₂)-Het, wherein 0<1<10, and R₂₁ & R₂₂, the same or different, are selected from the group of H, lower alkyl, and Het may be Het-1 or Het-2 or Het-3, or Het-4.

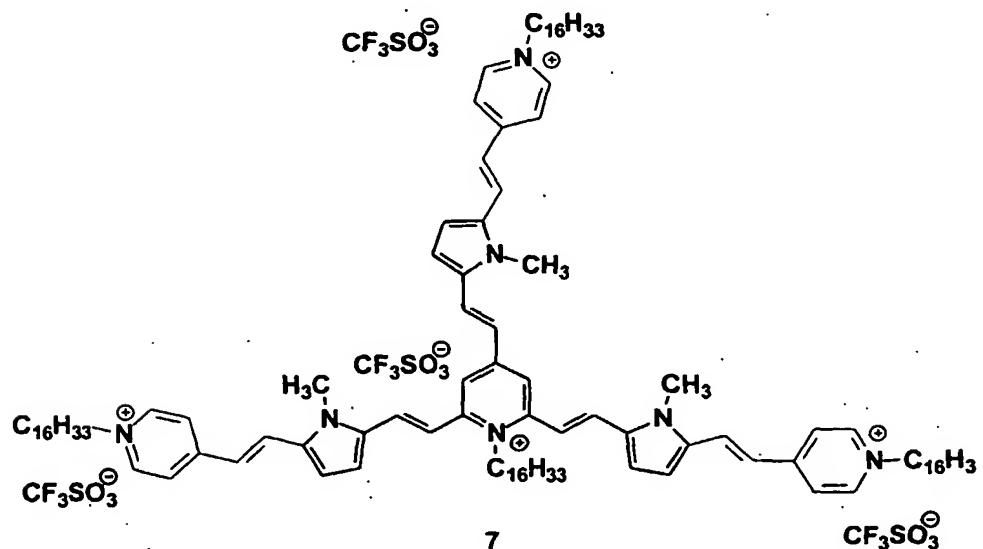
3. A compound according to claim 1, having the following formula (6)



4. A compound according to claim 2, having the following formula (3)



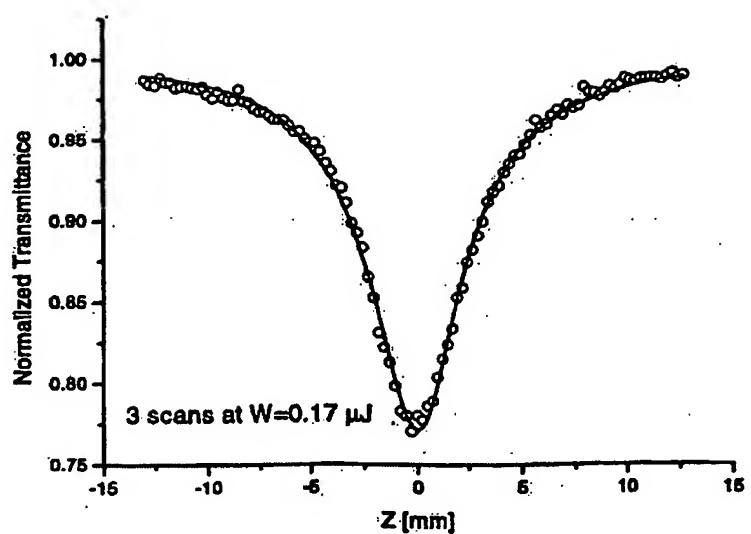
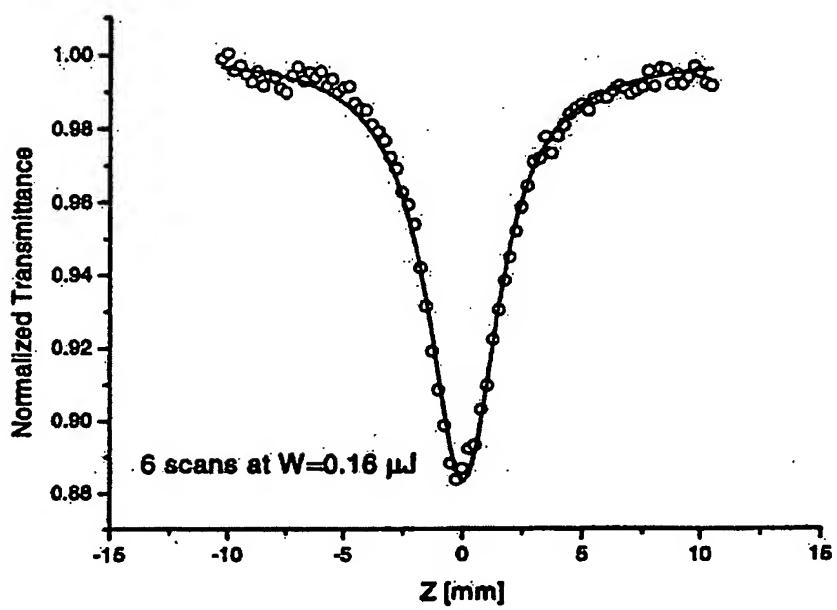
5. A compound according to claim 2, having the following formula (7)



10 6. Two-photon absorbing chromophore, in solution or in a solid state, characterized in that it is a compound of any of claim 1 to 5.

7. Compounds of general formula (I) according to claim 1 or (II) according to claim 2 for use in two-photon absorption systems.
8. Compounds of general formula (I) according to claim 1 or (II) according to claim 2 for use as optical power limiting agent via two-photon absorption.
- 5 9. Compounds of general formula (I) according to claim 1 or (II) according to claim 2 for use as imaging agents with two-photon absorbing activity for application in detection technologies such as two-photon laser scanning fluorescence microscopy.
10. A composition for use in two-photon absorption systems comprising a compound according to one of said claims 7, 8, 9.
- 10 11. A composition according to claim 10 characterized by the fact of comprising a polymer material chosen among poly(acrylate), poly(methacrylate), polyimide, polyamic acid, polystyrene, polycarbonate, polyurethane.
12. A composition according to claim 10 characterized by the fact of comprising an organically-modified silica (SiO_2) network.
- 15 13. A composition according to claim 10 and 11, characterized by the fact that the said compound is linked to the polymer materials by covalent bonds.
14. A composition according to claim 10 and 12, characterized by the fact that the said compound is linked to the silica network by covalent bonds.
- 20 15. A composition according to claim 10 for use as optical power limiting agent via two-photon absorption.
16. A composition according to claim 10 for use as imaging agent with two-photon absorbing activity for application in detection technologies such as two-photon laser scanning fluorescence microscopy.

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**Fig. 1****Fig. 2**

INTERNATIONAL SEARCH REPORT

International Application No

PCT/EP 07300

A. CLASSIFICATION OF SUBJECT MATTER

IPC 7 C07D401/14 C07D497/04 A61K41/00 A61K49/00
 //((C07D401/14, 213:00, 207:00, 207:00, 207:00), (C07D497/04, 333:00, 319:00))

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC 7 C07D A61K

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

EPO-Internal, CHEM ABS Data, MEDLINE

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	WO 01 70735 A (BOZIO RENATO ;PAGANI GIORGIO (IT); ABBOTTO ALESSANDRO (IT); UNI DE) 27 September 2001 (2001-09-27) cited in the application the whole document ---	1,2,6
X	ABBOTTO ALESSANDRO ET AL: "Novel heterocycle-based two-photon absorbing dyes." ORGANIC LETTERS. UNITED STATES 2 MAY 2002, vol. 4, no. 9, 2 May 2002 (2002-05-02), pages 1495-1498, XP002259759 ISSN: 1523-7060 page 1495, right-hand column -page 1496, right-hand column; table 1 --- -/-	1,6

Further documents are listed in the continuation of box C.

Patent family members are listed in annex.

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- *8* document member of the same patent family

Date of the actual completion of the international search

12 December 2003

Date of mailing of the international search report

29/12/2003

Name and mailing address of the ISA

European Patent Office, P.B. 5818 Patentlaan 2
 NL - 2280 HV Rijswijk
 Tel. (+31-70) 340-2040, Tx. 31 651 epo nl,
 Fax: (+31-70) 340-3016

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GOSS, I

INTERNATIONAL SEARCH REPORT

International Application No
PCT/EP 07300

C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

Category	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	KORNILOV M Y ET AL: "INTERACTION OF CHROMOPHORES IN SYMMETRICAL 2,5-SUBSTITUTED DERIVATIVES OF FURANE THIOPHENE" UKRAINIAN CHEMISTRY JOURNAL - UKRAINSKII KHIMICHESKII ZHURNAL, ALLERTON PRESS, NEW YORK, NY, US, vol. 40, no. 5, 1974, pages 52-54, XP001000456 ISSN: 1063-4568 tables I,II ---	1,6
X	KOSSMEHL G ET AL: "UEBER POLYARYLENALKENYLENE UND POLYHETEROARYLENALKENYLENE, 9 SYNTHESEN UND CHARAKTERISIERUNG VON POLY(5-METHYLTHIAZOL04,5-F BENZTHIAZOLIUM-2,6-DIYLVINYLENARYLENVINYLN)EN UND EINIGEN MODELLVERBINDUNGEN" MAKROMOLEKULARE CHEMIE, MACROMOLECULAR CHEMISTRY AND PHYSICS, HUTHIG UND WEPF VERLAG, BASEL, CH, vol. 180, no. 6, 1979, pages 1441-1463, XP001000460 ISSN: 0025-116X examples 9A-9F ---	1,6
A	BRADAMANTE S ET AL: "HETEROCYCLES AS DONOR AND ACCEPTOR UNITS IN PUSH-PULL CONJUGATED MOLECULES. PART I" JOURNAL OF PHYSICAL ORGANIC CHEMISTRY, WILEY, GB, vol. 10, no. 7, 1997, pages 514-524, XP001000454 ISSN: 0894-3230 the whole document -----	1,2,6

INTERNATIONAL SEARCH REPORT

Information on patent family members

International Application No

PCT/EP 017300

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		EP	1268467 A1	02-01-2003
		US	2003118916 A1	26-06-2003